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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO. CONFIRMATION N			
09/825,246	04/02/2001	Sharat Singh	0225-0033.20 4459			
33603	7590 01/20/2004		EXAMINER			
ACLARA BIOSCIENCES, INC. 1288 PEAR AVENUE			TUNG, JOYCE			
MOUNTAIN VIEW, CA 94043			ART UNIT	PAPER NUMBER		
			1637			
•			DATE MAILED: 01/20/2004	DATE MAILED: 01/20/2004		

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No	· ·	Applicant(s)					
Office Action Summary		09/825,246							
		Examiner		SINGH ET AL.  Art Unit					
		Joyce Tung		1637					
	The MAILING DATE of this communication app		er sheet with the c		idress				
THE	OF REPLY  HORTENED STATUTORY PERIOD FOR REPLY  MAILING DATE OF THIS COMMUNICATION.  Pensions of time may be available under the provisions of 37 CFR 1.13								
after - If the - If NO - Failu - Any	r SIX (6) MONTHS from the mailing date of this communication. e period for reply specified above is less than thirty (30) days, a reply D period for reply is specified above, the maximum statutory period w ure to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing	within the statutory n vill apply and will expir cause the application	ninimum of thirty (30) days re SIX (6) MONTHS from t n to become ABANDONEC	s will be considered timel the mailing date of this c	y. ommunication.				
Status	ed patent term adjustment. See 37 CFR 1.704(b).								
1)🛛	Responsive to communication(s) filed on 11 Se	eptember 2003.							
2a) <u></u> ☐	This action is <b>FINAL</b> . 2b)⊠ This action is non-final.								
3)[	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.								
Disposit	ion of Claims								
4)🖂	Claim(s) 16,17 and 19-29 is/are pending in the	application.							
·	4a) Of the above claim(s) is/are withdrawn from consideration.								
	Claim(s) is/are allowed.								
	Claim(s) <u>16-17 and 19-29</u> is/are rejected.								
	7) Claim(s) is/are objected to.								
	Claim(s) are subject to restriction and/or	election requir	ement.						
Applicat	ion Papers								
	The specification is objected to by the Examiner								
10)[_]	The drawing(s) filed on is/are: a) acce	•	•						
	Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).								
11)[]	Replacement drawing sheet(s) including the correction.		• • • •		` '				
	The oath or declaration is objected to by the Exa	ammer. Note tri	e attached Office /	Action or form P1	O-152.				
	Acknowledgment is made of a claim for ferring		\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	(1) (0)					
a)	Acknowledgment is made of a claim for foreign All b) Some * c) None of:  1. Certified copies of the priority documents			-(a) or (t).					
	2. Certified copies of the priority documents	have been rec	eived in Applicatio	on No					
	3. Copies of the certified copies of the priori	ty documents h	lave been received	d in this National	Stage				
* 5	application from the International Bureau See the attached detailed Office action for a list of			1					
13) 🗌 A	Acknowledgment is made of a claim for domestic	priority under	35 U.S.C. § 119(e)	) (to a provisional	application)				
3	ince a specific reference was included in the first 7 CFR 1.78.				Data Sheet.				
	)				a anacifia				
re	eference was included in the first sentence of the	specification of	or in an Application	Data Sheet. 37	CFR 1.78.				
					2				
Attachment	t(s) e of References Cited (PTO-892)		1						
	e of Draftsperson's Patent Drawing Review (PTO-948)		Interview Summary (F Notice of Informal Pa						
	nation Disclosure Statement(s) (PTO-1449) Paper No(s)		Other: .	,,					
D-1-1-1-	1.05								

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### **DETAILED ACTION**

## **Continued Prosecution Application**

- 1. The request filed on 9/11/2003 for a Continued Prosecution Application (CPA) under 37 CFR 1.53(d) based on parent Application No. 09/825,246 is acceptable and a CPA has been established. An action on the CPA follows.
- 2. Following the entry of the amendment filed 9/11/2003, the claims 16-17 and 19-29 are pending.
- 3. Applicant's arguments with respect to claims 16-17 and 19-29 have been considered but are most in view of the new ground(s) of rejection.

## Claim Rejections - 35 USC § 103

- 4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
  - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later

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invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

5. Claims 16-17, 19-21 and 23-28 are rejected under 35 U.S.C. 103(a) as being unpatentable over Grossman et al. (5,470,705) in view of Kline et al. (5,459,078).

Grossman et al. disclose a method of detecting a plurality of different sequences in a target sequence involving a plurality of sequence probes (See column 2, lines 54-56). The probe comprises the features of the e-tag probe as claimed in claims 16-17, 19-21 and 23-28. The probe includes a binding polymer, a polymer chain which imparts to that probe, a distinctive ratio of charge/translational frictional drag and a reporter attached to the binding polymer (See column 20, lines 52-57). The binding polymer is an oligonucleotide including at least 10-20 bases allowing hybridization to the target polynucleotide (See column 6, lines 66-67 and column 7, lines 1-10). Other binding polymers are analogs of polynucleotides, such as deoxynucleotides with thiophosphodiester linkage (See column 7, lines 11-19). The polymer chain has a ratio of charge/translational frictional drag which is evidenced by a distinctive electrophoretic mobility in a non-sieving matrix (See column 7, lines 50-64). The polymer chain can be polyethylene oxide (PEO) or a polypeptide chain where the chains are attached to different-sequence binding polymers (See column 3, lines 11-18). The teachings suggest that the charge/translational frictional drag is consisted of carbon, hydrogen, oxygen, phosphorus, nitrogen, sulfur and boron as recited in claim 24. The label refers to a fluorophore or chromophore (See column 6, lines 39-44). The features of Grossman et al.'s probe suggest the features of the claimed e-tag probe.

Grossman et al do not disclose the set of electrophoretic probe in which the oligonucleotide portion is attached with a capture ligand.

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Kline et al. disclose a competitive digoxin assay method (See the Abstract). A test sample suspected of containing the analyte of interest may be contacted with the capture reagent to form a charged capture reagent/analyte complex. The complex is then contacted to the oppositely charged solid phase to attract, attach and immobilize the capture reagent/analyte complex (See the Abstract). The test sample can be derived from any desired source (See column 8, lines 5-18). The analyte can be any substance for which there exists a naturally occurring specific binding member or for which a specific binding member can be prepared (See column 8, lines 19-32). The specific binding pair can be biotin and avidin, and complementary nucleotide sequences including probe and capture nucleic acid sequence used in DNA hybridization assays to detect a target nucleic acid sequence (See column 7, lines 37-53).

One of ordinary skill in the art would have been motivated to apply the binding pair biotin and avidin to the nucleic acid probe of Grossman to make the electrophoretic probes for detecting the presence of absence of one or more of a plurality nucleic sequence in a sample. Kline et al. disclose that the invention is not limited to immunoreactive assay and any assays using specific binding reactions between the analyte and assay reagents can be performed (See column 7, lines 28-33) and the ion-capture technique increases the potential number of complexes that can be immobilized on a solid support. It would have been <u>prima facie</u> obvious to make the set of electrophoretic probes with a capture ligand for detecting the presence or absence of one or more of a plurality nucleic acid sequence in a sample.

6. Claim 22 is rejected under 35 U.S.C. 103(a) as being unpatentable over Grossman et al. (5,470,705) in view of Kline et al. (5,459,078), as applied to claims 16-17, 19-21 and 23-28 above, and further in view of Huie et al. (5,470,967).

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The teachings of Grossman et al. and Kline et al. are set forth in section 5 above. None of the references discloses that said oligonucleotide has at least on nuclease resistant linkage

Huie et al. disclose phosphodiester linkage in oligonucleotide analogs (See column 3, lines 59-62) and phosphorothioate diester shows increased resistance to nuclease (See column 3, lines 59-67). Thus, it would have been <u>prima facie</u> obvious to one of ordinary skill in the art at the time of the instant invention to use phosphodiester linkage as indicated by Huie et al. in the oligonucleotide probe of Grossman et al. to resist nuclease activity because the use of modified linkage within the oligonucleotide makes them nuclease resistant (See column 3, lines 63-67).

7. Claim 29 is rejected under 35 U.S.C. 103(a) as being unpatentable over Grossman et al. (5,470,705 (1995)) in view of Kline et al. (5,459,078) as applied to claims above, and further in view of Ullman et al. (6,251,581B1 (2001))

The teachings of Grossman et al. and Kline et al. are set forth in section 5 above.

Grossman et al. and Kline et al. do not disclose the detectable labels, which are the compounds, listed in claim 29.

Ullman et al. disclose a method for determining an analyte in a medium (See the Abstract). The method applies a chemiluminescent compound associated with an specific binding pair member (See column 4, lines 54-65 and column 5, lines 8-14). The compound has the same structure as the compound listed in claims 29 (See column 42-58).

One of ordinary skill in the art at the time of the invention was made would have been motivated to apply the chemiluminescent compound of Ullman et al. to the probe of Grossman et al. in order to construct the set of electrophoretic tag probe of instant invention. Ullman et al. disclose a chemiluminescent compound to bind to a specific binding pair complex so that the

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detection may be performed without heating the medium to produce light and conducted at a constant temperature (See column 7, lines 28-31). If the analytes are proteins, by avoiding heating, protein analytes would not be inactivated and thus the sensitivity of the method is increased. It would have been <u>prima facie</u> obvious to apply the fluorescent molecules to the electrophoric release tag to construct the set of electrophoretic tag probe to avoid inactivating protein analytes. Thus it would have been <u>prima facie</u> obvious to apply the fluorescent molecules to the electrophoric release tag to construct the set of electrophoretic tag probe.

### Summary

- 8. No claims are allowable
- 9. Any inquiries concerning this communication or earlier communications from the examiner should be directed to Joyce Tung whose telephone number is (703) 305-7112. The examiner can normally be reached on Monday-Friday from 8:00 AM-4:30 PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached at (703) 308-1119 on Monday-Friday from 10:00 AM-6:00 PM.

Any inquiries of a general nature or relating to the status of this application should be directed to the Chemical/Matrix receptionist whose telephone number is (703) 308-0196.

10. Papers related to this application may be submitted to Group 1600 by facsimile transmission. Papers should be faxed to Art Unit 1637 via the PTO Fax Center located in Crystal Mall 1 using (703) 305-3014 or 308-4242. The faxing of such papers must conform with the notice published in the Official Gazette, 1096 OG 30 (November 15, 1989).

Joyce Tung 5.7 January 13, 2004

GARY BENZION, PH.D SUPERVISORY PATENT EXAMINER
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